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08/552,839 attribuent to Paper# 6

=> s adenovirus? or adenoviral

2421 ADENOVIRUS?

159 ADENOVIRAL

L1 2452 ADENOVIRUS? OR ADENOVIRAL

=> s l1 and late

25443 LATE

L2 1072 L1 AND LATE

=> s 12 and late gene?

25443 LATE

1274088 GENE?

120 LATE GENE?

(LATE (W) GENE?)

L3 46 L2 AND LATE GENE?

=> s 13 and inducib?

2972 INDUCIB?

L4 16 L3 AND INDUCIB?

=> d 14,1-16,cit,ab

1. 5,695,963, Dec. 9, 1997, Endothelial PAS domain protein; Steven L. McKnight, et al., 435/69.1, 7.1, 243, 325; 530/350; 536/23.5, 24.31 [IMAGE AVAILABLE]

US PAT NO: 5,695,963 [IMAGE AVAILABLE] L4: 1 of 16

ABSTRACT:

The invention provides methods and compositions relating to endothelial PAS domain protein 1 (EPAS1) and related nucleic acids. The proteins may be produced recombinantly from transformed host cells from the disclosed EPAS1 encoding nucleic acids or purified from human cells. The invention provides isolated EPAS1 hybridization probes and primers capable of specifically hybridizing with the disclosed EPAS1 gene, EPAS1-specific binding agents such as specific antibodies, and methods of making and using the subject compositions in diagnosis, therapy and in the biopharmaceutical industry.

2. 5,688,640, Nov. 18, 1997, Methods of screening of effectors of endothelin converting enzyme-1; Masashi Yanagisawa, 435/6, 24, 29, 69.1, 172.3, 252.3 [IMAGE AVAILABLE]

US PAT NO: 5,688,640 [IMAGE AVAILABLE] L4: 2 of 16

ABSTRACT:

Endothelin-1 (ET-1), pl-residue vasoactive peptide, produced in vascular endothelial cels from the 38-residue inactive intermediate, big endothelin-1 via a specific cleavage at Trp21-Val22. The protease that catalyzes the conversion, endothelin converting enzyme (ECE), constitutes a potential regulatory site for the production of the active peptide. Disclosed herein is the identification of ECE-1, a novel membrane-bound neutral Zn.sup.2+ metalloprotease that is expressed abundantly in endothelial cells in vivo, and structurally related to neutral endopeptidase 24.11 and Kell blood group protein. When transfected into cultured cells that normally secrete only big ET-1, the ECE-1 cDNA conferred the ability to secrete mature ET-1. In transfected cells, ECE-1 processes endogenously synthesized big ET-1 as well as exogenously supplied big ET-1, which interacts with ECE-1 on the cell surface. The remarkable specificity of ECE-1 provides a target for selective pharmacological intervention to alter ET-1 production in certain clinical disorders.

3. 5,672,479, Sep. 30, 1997, Methods for identifying compounds that bind to PUR protein; Edward M. Johnson, et al., 435/7.1, 7.23, 7.71, 7.93; 530/300, 358; 935/39, 41 [IMAGE AVAILABLE]

L4: 3 of 16

US PAT NO: 5,672,479 [IMAGE AVAILABLE]

ABSTRACT:

The present invention relates to the PUR protein, nucleotide sequences and expression vectors encoding PUR, and to methods for inhibiting PUR activity. Inhibitors of PUR activity may be used to treat hyperproliferative diseases such as cancer.

4. 5,665,590, Sep. 9, 1997, Method for isolating and directly cloning genes which encode cell-surface and secreted proteins; Zhi Yang, 435/6, 172.3, 320.1 [IMAGE AVAILABLE]

US PAT NO: 5,665,590 [IMAGE AVAILABLE] L4: 4 of 16

ABSTRACT:

Methods and DNA cloning vectors are provided for obtaining genes encoding secreted proteins and cell surface proteins.

5. 5,658,761, Aug. 19, 1997, Stromal cell lines from human bone marrow and their use; Karin Thalmeier, et al., 435/69.4, 69.51, 69.52, 172.1, 172.3, 372, 373 [IMAGE AVAILABLE]

US PAT NO: 5,658,761 [IMAGE AVAILABLE] L4: 5 of 16

ABSTRACT:

A human bone marrow stromal cell line, which is characterized in that the cells of the cell line, after irradiation which results in the growth being arrested, remain adherent, is suitable for use as a feeder layer for supporting the proliferation of blood cells.

6. 5,652,224, Jul. 29, 1997, Methods and compositions for gene therapy for the treatment of defects in lipoprotein metabolism; James M. Wilson, et al., 514/44; 424/93.21; 435/172.3, 320.1, 325, 354, 366, 369, 370 [IMAGE AVAILABLE]

US PAT NO: 5,652,224 [IMAGE AVAILABLE] L4: 6 of 16

ABSTRACT:

The invention provides a recombinant viral vector comprising the DNA of, or corresponding to, at least a portion of the genome of an adenovirus, which portion is capable of infecting a hepatic cell; and a human VLDL receptor gene operatively linked to regulatory sequences directing its expression. The vector is capable of expressing the normal

VLDL receptor gene product in hepatic cells in vivo or in vitro. This viral vector is useful in the treatment of metabolic derders caused by the accumulation of LDL in plasma, such as familial hypercholesterolemia or familial combined hyperlipidemia.

7. 5,650,550, Jul. 22, 1997, Mutant mice having a deficit of functional estrogen receptors; Kenneth S. Korach, et al., 800/2; 435/172.3, 354; 800/DIG.1; 935/10, 70 [IMAGE AVAILABLE]

US PAT NO: 5,650,550 [IMAGE AVAILABLE] L4: 7 of 16

ABSTRACT:

The present invention provides a mutant non-human vertebrate, in which all or some of the germ and somatic cells contain a mutation in at least one steroid hormone receptor allele, which mutation is introduced into the vertebrate, or an ancestor of the vertebrate, at an embryonic stage, and which mutation produces a phenotype in the vertebrate characterized by a deficit of functional steroid hormone receptors encoded by the allele. Also disclosed are related methods and constructs.

8. 5,641,670, Jun. 24, 1997, Protein production and protein delivery; Douglas A. Treco, et al., 435/254.11, 320.1 [IMAGE AVAILABLE]

US PAT NO: 5,641,670 [IMAGE AVAILABLE] L4: 8 of 16

ABSTRACT:

The invention relates to constructs comprising: a) a targeting sequence; b) a regulatory sequence; c) an exon; and d) an unpaired splice-donor site. The invention further relates to a method of producing protein in vitro or in vivo comprising the homologous recombination of a construct as described above within a cell. The homologously recombinant cell is then maintained under conditions which will permit transcription and translation, resulting in protein expression. The present invention further relates to homologously recombinant cells, including primary, secondary, or immortalized vertebrate cells, methods of making the cells, methods of homologous recombination to produce fusion genes, methods of altering gene expression in the cells, and methods of making a protein in a cell employing the constructs of the invention.

9. 5,639,661, Jun. 17, 1997, Genes and proteins for treating cystic fibrosis; Michael J. Welsh, et al., 435/252.3, 320.1; 536/23.5, 24.3 [IMAGE AVAILABLE]

US PAT NO: 5,639,661 [IMAGE AVAILABLE] L4: 9 of 16

ABSTRACT:

Disclosed are genes encoding novel CF monomer proteins which have cystic fibrosis transmembrane conductance regulator (CFTR) protein function.

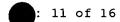
10. 5,627,033, May 6, 1997, Mammalian expression vectors; John M. Smith, et al., 435/6, 91.41, 172.3, 320.1, 325, 358, 365 [IMAGE AVAILABLE]

US PAT NO: 5,627,033 [IMAGE AVAILABLE] L4: 10 of 16

ABSTRACT:

A vector system that allows the rapid and effective screening of recombinant constructs. The vector system includes a marker protein useful for identifying transfected cell lines, wherein the promoter used to express the marker protein has be substantially weakened in comparison to its corresponding wild type form.

11. 5,589,358, Dec. 31, 1996, Ileal bile acid transporter compositions and methods; Paul A. Dawson, 435/69.1, 172.3, 252.3, 320.1, 325, 348, 350, 352, 358, 364, 365, 367, 369; 530/846; 536/23.5, 24.3 [IMAGE AVAILABLE]



ABSTRACT:

Bile acids are absorbed from the small intestine by an passive and an active mechanism. The active mechanism involves a Na.sup.+ /bile acid cotransporter. A cDNA encoding the ileal bile acid cotransporter has been isolated and sequenced. The amino acid sequence of the cotransporter protein is also disclosed. The renal bile acid cotransporter is also shown to be identical to the ileal cotransporter. The cotransporter disclosed herein will have use in treatment of hypercholesterolemia, diabetes, heart disease and liver disease. In addition, methods of screening man made and naturally occurring substances for the discovery of new bile acid transport inhibitors and activators and methods of detecting mutations in human ileal/renal bile acid transporter genes are disclosed.

12. 5,532,339, Jul. 2, 1996, Fusion protein between human macif and a heterologous pi anchor domain; Motowo Tomita, et al., 530/350; 435/69.7; 530/324, 380 [IMAGE AVAILABLE]

US PAT NO:

5,532,339 [IMAGE AVAILABLE]

L4: 12 of 16

ABSTRACT:

Fusion proteins comprising the extracellular domain of the human MACIF (Membrane Attack Complex Inhibition Factor) gene product and a heterologous phosphatidylinositol (PI) anchor domain are provided.

5,514,578, May 7, 1996, Polynucleotides encoding insect steroid hormone receptor polypeptides and cells transformed with same; David S. Hogness, et al., 435/325, 252.3, 348; 536/23.5 [IMAGE AVAILABLE]

US PAT NO:

5,514,578 [IMAGE AVAILABLE]

L4: 13 of 16

ABSTRACT:

Polynucleotide sequences which encode ecdysone receptors have been isolated and expressed in host cells.

14. 5,474,935, Dec. 12, 1995, Adeno-associated virus (AAV)-based eucaryotic vectors; Saswati Chatterjee, et al., 435/320.1; 424/93.1, 93.2; 435/172.3; 935/22, 32, 57 [IMAGE AVAILABLE]

US PAT NO:

5,474,935 [IMAGE AVAILABLE]

L4: 14 of 16

ABSTRACT:

The present invention relates to adeno-associated virus (AAV)-based eucaryotic vectors and uses thereof. Such vectors may, for example, be used to down regulate any targeted viral or cellular gene whose sequence is known. Furthermore, the vectors may also be used to cause the expression of proteins.

15. 5,428,070, Jun. 27, 1995, Treatment of vascular degenerative diseases by modulation of endogenous nitric oxide production of activity; John P. Cooke, et al., 514/557, 310 [IMAGE AVAILABLE]

US PAT NO:

5,428,070 [IMAGE AVAILABLE]

L4: 15 of 16

ABSTRACT:

Atherogenesis and restenosis are treated by long term administration of physiologically acceptable compounds which enhance the level of endogenous nitric oxide in the host. Alternatively, or in combination, other compounds may be administered which provide for short term enhancement of nitric oxide, either directly or by physiological processes. In addition, cells may be genetically engineered to provide a component in the synthetic pathway to nitric oxide, so as drive the

process to enhance nitric oxide concentration, particularly in conjunction with the inistration of a nitric oxide cursor

16. 4,988,624, Jan. 29, 1991, Lymphotoxin DNA, lymphotoxin expression vector; Tetsu Kakutani, et al., 435/320.1, 172.3; 536/23.2, 23.5, 23.51, 24.1; 935/27, 36 [IMAGE AVAILABLE]

L4: 16 of 16

US PAT NO: 4,988,624 [IMAGE AVAILABLE]

55 PAI NO. 4,500,024 [IMAGE AVAILABLE]

ABSTRACT:

A chromosomal DNA sequence which codes for human lymphotoxin, a lymphotoxin expression vector which contains a DNA sequence wherein a chromosomal DNA sequence coding for human lymphotoxin and promoter region which functions in animal cell are linked together, lymphotoxin resistant cell line, transformed animal cell culture which is formed by transforming cultured animal cell with a lymphotoxin expression vector which contains a chromosomal DNA sequence coding for human lymphotoxin and, a process for preparing human lymphotoxin, which comprises transforming cultured animal cell with a lymphotoxin expression vector which contains a chromosomal DNA sequence coding for human lymphotoxin, culturing the transformed cell culture to produce human lymphotoxin, and collecting the human lymphotoxin.

According to the present invention, LT which is expected for application as the antitumor agent can be produced effectively in a large amount.

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? begin 5,6,55,154,155,156,312,399,351,biotech,biosci

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(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: YL010 No. References: 84
16219119
Title: The tripartite leader sequence of subgroup C adenovirus major
    late mRNAs can increase the efficiency of mRNA export
Author(s): Huang W; Flint SJ (REPRINT)
Corporate Source: PRINCETON UNIV, DEPT MOL BIOL/PRINCETON//NJ/08544
    (REPRINT); PRINCETON UNIV, DEPT MOL BIOL/PRINCETON//NJ/08544
Journal: JOURNAL OF VIROLOGY, 1998, V72, N1 (JAN), P225-235
                  Publication date: 19980100
ISSN: 0022-538X
Publisher: AMER SOC MICROBIOLOGY, 1325 MASSACHUSETTS AVENUE, NW,
    WASHINGTON, DC 20005-4171
                                              (ABSTRACT AVAILABLE)
Language: English
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14779058
           Genuine Article#: UL104
                                     No. References: 91
Title: MESSENGER-RNA EXPORT CORRELATES WITH ACTIVATION OF TRANSCRIPTION IN
    HUMAN SUBGROUP-C ADENOVIRUS-INFECTED CELLS
Author(s): YANG UC; HUANG W; FLINT SJ
Corporate Source: PRINCETON UNIV, DEPT MOLEC BIOL/PRINCETON//NJ/08544;
    PRINCETON UNIV, DEPT MOLEC BIOL/PRINCETON//NJ/08544
Journal: JOURNAL OF VIROLOGY, 1996, V70, N6 (JUN), P4071-4080
ISSN: 0022-538X
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DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: TN847
                                     No. References: 50
Title: DEVELOPMENT OF CELL-LINES CAPABLE OF COMPLEMENTING E1, E4, AND
    PROTEIN-IX DEFECTIVE ADENOVIRUS TYPE-5 MUTANTS
Author(s): KROUGLIAK V; GRAHAM FL
Corporate Source: MCMASTER UNIV, DEPT BIOL, 1280 MAIN ST W/HAMILTON/ON L8S
    4K1/CANADA/; MCMASTER UNIV, DEPT BIOL/HAMILTON/ON L8S 4K1/CANADA/;
    MCMASTER UNIV, DEPT PATHOL/HAMILTON/ON L8S 4K1/CANADA/
Journal: HUMAN GENE THERAPY, 1995, V6, N12 (DEC), P1575-1586
ISSN: 1043-0342
Language: ENGLISH
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(c) 1998 Inst for Sci Info. All rts. reserv.
14402200
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Title: EFFICIENT DUAL TRANSCOMPLEMENTATION OF ADENOVIRUS E1 AND E4
    REGIONS FROM A 293-DERIVED CELL-LINE EXPRESSING A MINIMAL E4 FUNCTIONAL
    UNIT
Author(s): YEH P; DEDIEU JF; ORSINI C; VIGNE E; DENEFLE P; PERRICAUDET M
Corporate Source: RHONE POULENC RORER GENCELL, INST GUSTAVE ROUSSY, VIRUS
    ONCOGENES LAB, CNRS, URA 1301/F-94805 VILLEJUIF//FRANCE/; RHONE POULENC
    RORER GENCELL, CTR RECH VITRY ALFORTVILLE/F-94403 VITRY//FRANCE/
Journal: JOURNAL OF VIROLOGY, 1996, V70, N1 (JAN), P559-565
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Title: IDENTIFICATION OF A HIGH-MOLECULAR-WEIGHT CELLULAR PROTEIN COMPLEX
    CONTAINING THE ADENOVIRUS DNA-BINDING PROTEIN
Author(s): RICIGLIANO JW; BROUGH DE; KLESSIG DF
Corporate Source: RUTGERS STATE UNIV, WAKSMAN INST MICROBIOL, POB
    759/PISCATAWAY//NJ/08855; RUTGERS STATE UNIV, WAKSMAN INST
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Journal: VIROLOGY, 1994, V202, N2 (AUG 1), P715-723

ISSN: 0042-6822
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12689995 Genuine Article#: MF888 No. References: 77
Title: AN AP-1 BINDING-SITE IN THE UPSTREAM REGION OF DEB-A IS PART OF A

DEVELOPMENTALLY-REGULATED NEGATIVE ELEMENT

Author(s): WANG GL; GOLDSTEIN ES

Corporate Source: ARIZONA STATE UNIV, DEPT ZOOL/TEMPE//AZ/85287; ARIZONA STATE UNIV, DEPT ZOOL/TEMPE//AZ/85287

Journal: BIOCHIMICA ET BIOPHYSICA ACTA, 1993, V1216, N1 (OCT 19), P94-104

ISSN: 0006-3002

Language: ENGLISH Document Type: ARTICLE (Abstract Available)

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12507596 Genuine Article#: LR462 No. References: 22
Title: ENHANCED GEL MOBILITY SHIFT ASSAY FOR DNA-BINDING FACTORS
Author(s): HASSANAIN HH; DAI W; GUPTA SL
Corporate Source: HIPPLE CANC RES CTR,4100 S KETTERING

BLVD/DAYTON//OH/45439; HIPPLE CANC RES CTR,4100 S KETTERING

BLVD/DAYTON//OH/45439

Journal: ANALYTICAL BIOCHEMISTRY, 1993, V213, N1 (AUG 15), P162-167

ISSN: 0003-2697

Language: ENGLISH Document Type: ARTICLE

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Display 3/3/8 (Item 8 from file: 434) DIALOG(R)File 434:Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv.

12328766 Genuine Article#: LC210 No. References: 52

Title: EFFECT OF THE NONCONSERVED N-TERMINUS ON THE DNA-BINDING ACTIVITY OF THE YEAST TATA-BINDING PROTEIN

Author(s): KUDDUS R; SCHMIDT MC

Corporate Source: UNIV PITTSBURGH, SCH MED, DEPT MOLEC GENET &

BIOCHEM/PITTSBURGH//PA/15261; UNIV PITTSBURGH, SCH MED, DEPT MOLEC GENET & BIOCHEM/PITTSBURGH//PA/15261

Journal: NUCLEIC ACIDS RESEARCH, 1993, V21, N8 (APR 25), P1789-1796

ISSN: 0305-1048

Language: ENGLISH Document Type: ARTICLE (Abstract Available)

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12327416 Genuine Article#: LC650 No. References: 38

Title: STRUCTURE AND EXPRESSION IN ESCHERICHIA-COLI OF THE GENE CODING FOR PROTEIN P10 OF AFRICAN SWINE FEVER VIRUS

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Author(s): MUNOZ M; FREIJE JMP; SALAS ML; VINUELA E; LOPEZOTIN C
Corporate Source: UNI UTONOMA MADRID, FAC CIENCIAS, CS CTR
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Title: A NOVEL NEGATIVE ELEMENT IN THE PROMOTER OF THE MOUSE
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Author(s): LIN ZH; EDENBERG HJ; CARR LG
Corporate Source: INDIANA UNIV, SCH MED, DEPT MED, 975 W WALNUT
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Journal: JOURNAL OF BIOLOGICAL CHEMISTRY, 1993, V268, N14 (MAY 15), P
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Title: 3 DISTINCT NUCLEAR-PROTEIN BINDING-SITES IN THE PROMOTER OF THE
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Author(s): YU LJ; COHEN D; PIEKARZ RL; HORWITZ SB
Corporate Source: YESHIVA UNIV ALBERT EINSTEIN COLL MED, DEPT MOLEC
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Journal: JOURNAL OF BIOLOGICAL CHEMISTRY, 1993, V268, N10 (APR 5), P
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(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: KR218
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12159486
Title: JUNB DIFFERS FROM C-JUN IN ITS DNA-BINDING AND DIMERIZATION DOMAINS,
    AND REPRESSES C-JUN BY FORMATION OF INACTIVE HETERODIMERS
Author(s): DENG TL; KARIN M
Corporate Source: UNIV CALIF SAN DIEGO, SCH MED, CTR MOLEC GENET, DEPT
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    GENET, DEPT PHARMACOL/LA JOLLA//CA/92093
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Journal: GENES & DEVELOPMENT, 1993, V7, N3 (MAR), P479-490 ISSN: 0890-9369 Document Type: ARTICLE (Abstract Available) Language: ENGLISH - end of record -? (Item 13 from file: 434) Display 3/3/13 DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: GA643 No. References: 63 11028472 Title: PROTEINS OF THE NUCLEAR FACTOR-I FAMILY ACT AS AN ACTIVATOR OF THE LATE PROMOTER IN HUMAN POLYOMAVIRUS BK INVITRO Author(s): CHAKRABORTY T; DAS GC Corporate Source: UNIV TEXAS, HLTH SCI CTR, DEPT MOLEC BIOL, POB 2003/TYLER//TX/75710; UNIV TEXAS, HLTH SCI CTR, DEPT MOLEC BIOL, POB 2003/TYLER//TX/75710 Journal: JOURNAL OF GENERAL VIROLOGY, 1991, V72, AUG, P1935-1942 (Abstract Available) Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/14 (Item 14 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: FT820 No. References: 27 10939966 Title: CONTROL OF TRANSCRIPTION INVITRO FROM SIMIAN VIRUS-40 PROMOTERS BY PROTEINS FROM VIRAL MINICHROMOSOMES Author(s): BEARD P; BRUGGMANN H Corporate Source: SWISS INST EXPTL CANC RES/CH-1066 EPALINGES//SWITZERLAND/ Journal: CURRENT TOPICS IN MICROBIOLOGY AND IMMUNOLOGY, 1989, V144, P47-54 Language: ENGLISH Document Type: REVIEW - end of record -? Display 3/3/15 (Item 15 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: FB496 10717711 No. References: 31 Title: CHARACTERIZATION OF SV40 ENHANCER MOTIFS INVOLVED IN POSITIVE AND NEGATIVE REGULATION OF THE CONSTITUTIVE LATE PROMOTER ACTIVITY - EFFECT OF T-ANTIGEN Author(s): SCIELLER P; OMILLI F; BORDE J; MAY E Corporate Source: INST RECH SCI CANC, ONCOL MOLEC LAB/F-94800 VILLEJUIF//FRANCE/; INST RECH SCI CANC, ONCOL MOLEC LAB/F-94800 VILLEJUIF//FRANCE/ Journal: VIROLOGY, 1991, V181, N2, P783-786 Language: ENGLISH Document Type: NOTE - end of record -? Display 3/3/16 (Item 16 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: R3666 No. References: 39 09193652 Title: HUMAN ETHANOL-INDUCIBLE P450IIE1 - COMPLETE GENE SEQUENCE,

PROMOTER CHARACTERIZATION, CHROMOSOME MAPPING, AND CDNA-DIRECTED

EXPRESSION

Author(s): UMENO M; MCBRIDE OW; YANG CS; GELBOIN HV; GONZALEZ FJ Corporate Source: NCI LEC CARCINOGENESIS LAB/BETHESD MD/20892; NCI, BIOCHEM LAB/BETHESDA//MD/20892; UNIV MED & DEN. NEW JERSEY, NEW JERSEY MED SCH, DEPT BIOCHEM/NEWARK//NJ/07103 Journal: BIOCHEMISTRY, 1988, V27, N25, P9006-9013 Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/17 (Item 17 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. 09110376 Genuine Article#: Q6869 No. References: 49 Title: CONCENTRATION-DEPENDENCE OF TRANSCRIPTIONAL TRANSACTIVATION IN INDUCIBLE E1A-CONTAINING HUMAN-CELLS Author(s): BRUNET LJ; BERK AJ Corporate Source: UNIV CALIF LOS ANGELES, DEPT MICROBIOL/LOS ANGELES//CA/90024; UNIV CALIF LOS ANGELES, DEPT MOLEC BIOL/LOS ANGELES//CA/90024 Journal: MOLECULAR AND CELLULAR BIOLOGY, 1988, V8, N11, P4799-4807 Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/18 (Item 18 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. 09089549 Genuine Article#: Q5807 No. References: 34 Title: TARGETING OF AN INDUCIBLE TOXIC PHENOTYPE IN ANIMAL-CELLS Author(s): BORRELLI E; HEYMAN R; HSI M; EVANS RM Corporate Source: SALK INST BIOL STUDIES, HOWARD HUGHES MED INST, GENE EXPRESS LAB/LA JOLLA//CA/92037 Journal: PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA, 1988, V85, N20, P7572-7576 Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/19 (Item 19 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: P3983 No. References: 46 08923273 Title: EXCESS ANTISENSE RNA FROM INFECTIOUS RECOMBINANT SV40 FAILS TO INHIBIT EXPRESSION OF A TRANSFECTED, INTERFERON-INDUCIBLE GENE Author(s): KERR SM; STARK GR; KERR IM Corporate Source: IMPERIAL CANC RES FUND LABS, POB 123, LINCOLNS INN FIELDS/LONDON WC2A 3PX//ENGLAND/ Journal: EUROPEAN JOURNAL OF BIOCHEMISTRY, 1988, V175, N1, P65-73 Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/20 (Item 20 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. 08871590 Genuine Article#: P0741 No. References: 69

Title: MOLECULAR CHARACTERIZATION AND CHROMOSOMAL MAPPING OF MELANOMA

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GROWTH STIMULATORY ACTIVITY, A GROWTH-FACTOR STRUCTURALLY RELATED TO
    BETA-THROMBOGLOBU
Author(s): RICHMOND A; BALENTIEN E; THOMAS HG; FLAGGS G; BARTON DE; SPIESS
    J; BORDONI R; FRANCKE U; DERYNCK R
Corporate Source: VET ADM MED CTR/ATLANTA//GA/00000; EMORY UNIV, SCH
    MED/ATLANTA//GA/30322; GENENTECH INC, DEPT MOLEC BIOL/S SAN
    FRANCISCO//CA/94080; YALE UNIV, SCH MED, DEPT HUMAN GENET/NEW
    HAVEN//CT/06510; SALK INST BIOL STUDIES, MAX PLANCK RES PROGRAM/SAN
    DIEGO//CA/92138
Journal: EMBO JOURNAL, 1988, V7, N7, P2025-2033
Language: ENGLISH Document Type: ARTICLE
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                         (Item 21 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: N6338 No. References: 52
08826865
Title: UPSTREAM REGULATORY REGION FOR INDUCIBLE EXPRESSION OF THE
    CHICKEN SKELETAL MYOSIN ALKALI LIGHT-CHAIN GENE
Author(s): SHIRAKATA M; NABESHIMA YI; KONISHI K; FUJIIKURIYAMA Y
Corporate Source: JAPANESE FDN CANC RES, INST CANC, DEPT BIOCHEM/TOKYO
    170//JAPAN/; TOHOKU UNIV, FAC SCI, INST BIOL/SENDAI/MIYAGI 980/JAPAN/
Journal: MOLECULAR AND CELLULAR BIOLOGY, 1988, V8, N6, P2581-2588
Language: ENGLISH Document Type: ARTICLE
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                         (Item 22 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
         Genuine Article#: N2825 No. References: 39
08768339
Title: ENHANCER-LIKE INTERFERON RESPONSIVE SEQUENCES OF THE HUMAN AND
    MURINE (2'-5') OLIGOADENYLATE SYNTHETASE GENE PROMOTERS
Author(s): COHEN B; PERETZ D; VAIMAN D; BENECH P; CHEBATH J
Corporate Source: WEIZMANN INST SCI, DEPT VIROL/IL-76100 REHOVOT//ISRAEL/
Journal: EMBO JOURNAL, 1988, V7, N5, P1411-1419
Language: ENGLISH Document Type: ARTICLE
                                 - end of record -
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                         (Item 23 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: M3557
                                    No. References: 24
Title: PRODUCTION OF RECOMBINANT HUMAN CSF-1 IN AN INDUCIBLE
    MAMMALIAN EXPRESSION SYSTEM
Author(s): WEAVER JF; MCCORMICK F; MANOS MM
Corporate Source: CETUS CORP, DEPT ANALYT DEV, 1400 53RD
    ST/EMERYVILLE//CA/94608; CETUS CORP, DEPT MOLEC
    BIOL/EMERYVILLE//CA/94608
Journal: BIO-TECHNOLOGY, 1988, V6, N3, P287-290
Language: ENGLISH Document Type: ARTICLE
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                         (Item 24 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
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(c) 1998 Inst for Sci Info. All rts. reserv.
          Genuine Article#: L7672
                                     No. References: 2
Title: ESTABLISHMENT OF A TEMPERATURE-INDUCIBLE CELL-LINE FOR
    HUMAN-PLASMINOGEN ACTIVATOR (TISSUE-TYPE) BY TRANSFECTION OF MONKEY
    CELLS WITH EXPRESSION CONSTRUCTS
Author(s): WEIDLE UH; LAWETZKY A; BUCKEL P
Corporate Source: BOEHRINGER MANNHEIM GMBH, BIOCHEM RES CTR, BAHNHOFSTR
    9-15/D-8132 TUTZING//FED REP GER/
Journal: GENE, 1987, V59, N2-3, P231-239
Language: ENGLISH Document Type: ARTICLE
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      Display 3/3/25
                         (Item 25 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: J9942
08325127
                                     No. References: 41
Title: IDENTIFICATION AND CHARACTERIZATION OF THE HUMAN CYTOMEGALO-VIRUS
    IMMEDIATE-EARLY REGION-2 GENE THAT STIMULATES GENE-EXPRESSION FROM AN
    INDUCIBLE PROMOTER
Author(s): HERMISTON TW; MALONE CL; WITTE PR; STINSKI MF
Corporate Source: UNIV IOWA, DEPT MICROBIOL/IOWA CITY//IA/52242
Journal: JOURNAL OF VIROLOGY, 1987, V61, N10, P3214-3221
Language: ENGLISH Document Type: ARTICLE
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                         (Item 26 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
          Genuine Article#: J4730 No. References: 57
08239919
Title: AN INDUCIBLE MAMMALIAN AMBER SUPPRESSOR - PROPAGATION OF A
    POLIOVIRUS MUTANT
Author(s): SEDIVY JM; CAPONE JP; RAJBHANDARY UL; SHARP PA
Corporate Source: MIT, CTR CANC RES/CAMBRIDGE//MA/02139; MIT, DEPT
    BIOL/CAMBRIDGE//MA/02139
Journal: CELL, 1987, V50, N3, P379-389
Language: ENGLISH Document Type: ARTICLE
                                 - end of record -
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      Display 3/3/27
                         (Item 27 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: H2971
                                     No. References: 39
08084270
Title: INTERFERON-BETA GENE-REGULATION - TANDEMLY REPEATED SEQUENCES OF A
    SYNTHETIC 6-BP OLIGOMER FUNCTION AS A VIRUS-INDUCIBLE ENHANCER
Author(s): FUJITA T; SHIBUYA H; HOTTA H; YAMANISHI K; TANIGUCHI T
Corporate Source: OSAKA UNIV, INST MOLEC & CELLULAR
    BIOL, YAMADAOKA1-3/SUITA/OSAKA 565/JAPAN/
Journal: CELL, 1987, V49, N3, P357-367
Language: ENGLISH Document Type: ARTICLE
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                         (Item 28 from file: 434)
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DIALOG(R) File 434: Scisearch(R) Cited Ref Sci

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          Genuine Article#: G3034 No. References: 6
07927106
Title: THE INDUCIBLE IAC OPERATOR-REPRESSOR SYSTEM IS FUNCTIONAL IN
    MAMMALIAN-CELLS
Author(s): HU MCT; DAVIDSON N
Corporate Source: CALTECH, DIV CHEM/PASADENA//CA/91125
Journal: CELL, 1987, V48, N4, P555-566
Language: ENGLISH Document Type: ARTICLE
                                 - end of record -
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      Display 3/3/29
                         (Item 29 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
         Genuine Article#: G1277 No. References: 42
07901497
Title: MUTATION OF THE C-FOS GENE DYAD SYMMETRY ELEMENT INHIBITS SERUM
    INDUCIBILITY OF TRANSCRIPTION INVIVO AND THE NUCLEAR REGULATORY
    FACTOR BINDING INVITRO
Author(s): GREENBERG ME; SIEGFRIED Z; ZIFF EB
Corporate Source: NYU MED CTR, DEPT BIOCHEM/NEW YORK//NY/10016; NYU MED
    CTR, KAPLAN CANC CTR/NEW YORK//NY/10016
Journal: MOLECULAR AND CELLULAR BIOLOGY, 1987, V7, N3, P1217-1225
Language: ENGLISH Document Type: ARTICLE
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                       (Item 30 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
          Genuine Article#: F9763
                                   No. References: 43
Title: LOSS OF (2'-5')OLIGOADENYLATE SYNTHETASE-ACTIVITY BY PRODUCTION OF
    ANTISENSE RNA RESULTS IN LACK OF PROTECTION BY INTERFERON FROM
    VIRAL-INFECTIONS
Author(s): DEBENEDETTI A; PYTEL BA; BAGLIONI C
Corporate Source: SUNY ALBANY, DEPT BIOL SCI/ALBANY//NY/12222
Journal: PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED
    STATES OF AMERICA, 1987, V84, N3, P658-662
Language: ENGLISH Document Type: ARTICLE
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      Display 3/3/31
                       (Item 31 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: E9568
07654678
                                    No. References: 69
Title: MULTIPLE HORMONE-INDUCIBLE ENHANCERS AS MEDIATORS OF
    DIFFERENTIAL TRANSCRIPTION
Author(s): TOOHEY MG; MORLEY KL; PETERSON DO
Corporate Source: TEXAS A&M UNIV, TEXAS AGR EXPT STN, DEPT BIOCHEM &
    BIOPHYS/COLLEGE STN//TX/77843
Journal: MOLECULAR AND CELLULAR BIOLOGY, 1986, V6, N12, P4526-4538
Language: ENGLISH Document Type: ARTICLE
                                 - end of record -
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                         (Item 32 from file: 434)
      Display 3/3/32
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DIALOG(R) File 434: Scisearch(R) Cited Ref Sci

(c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: D8063 No. References: 2 Title: AN INDUCIBLE EUKARYOTIC HOST-VECTOR EXPRESSION SYSTEM -AMPLIFICATION OF GENES UNDER THE CONTROL OF THE POLYOMA LATE PROMOTER IN A CELL-LINE PRODUCING A THERMOLABILE LARGE T-ANTIGEN Author(s): KERN FG; BASILICO C Corporate Source: NYU, SCH MED, DEPT PATHOL/NEW YORK//NY/10016 Journal: GENE, 1986, V43, N3, P237-245 Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/33 (Item 33 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: C9492 No. References: 29 07379486 Title: EFFECT OF E1A AND E1B VIRAL-PROTEINS ON THE EXPRESSION OF A CALCIUM IONOPHORE-INDUCIBLE GENE AND ITS PROMOTER Author(s): LIN AY; LEE AS Corporate Source: UNIV SO CALIF, SCH MED, DEPT BIOCHEM/LOS ANGELES//CA/90033; UNIV SO CALIF, SCH MED, CTR COMPREHENS CANC/LOS ANGELES//CA/90033 Journal: NUCLEIC ACIDS RESEARCH, 1986, V14, N12, P4911-4921 Language: ENGLISH Document Type: ARTICLE - end of record -? Display 3/3/34 (Item 34 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: A5751 No. References: 37 07176022 Title: A CALCIUM IONOPHORE-INDUCIBLE CELLULAR PROMOTER IS HIGHLY-ACTIVE AND HAS ENHANCERLIKE PROPERTIES Author(s): LIN AY; CHANG SC; LEE AS Corporate Source: UNIV SO CALIF, SCH MED, DEPT BIOCHEM/LOS ANGELES//CA/90033 Journal: MOLECULAR AND CELLULAR BIOLOGY, 1986, V6, N4, P1235-1243 Language: ENGLISH Document Type: ARTICLE - end of display -?

FILE 'USPAT' ENTERED AT 16:01:24 ON 11 FEB 1998

* WELCOME TO THE *

* U.S. PATENT TEXT FILE *

=> s adenovir? and E2A and promoter?

2484 ADENOVIR?

161 E2A

28681 PROMOTER?

L1 19 ADENOVIR? AND E2A AND PROMOTER?

=> s l1 and inducible

2870 INDUCIBLE

L2 10 L1 AND INDUCIBLE

=> s 12 and inducible promoter? (10A) E2A

2870 INDUCIBLE

28681 PROMOTER?

952 INDUCIBLE PROMOTER?

(INDUCIBLE (W) PROMOTER?)

161 E2A

0 INDUCIBLE PROMOTER? (10A) E2A

L3 0 L2 AND INDUCIBLE PROMOTER? (10A) E2A

=> d 12, 1-10, cit, ab

1. 5,695,995, Dec. 9, 1997, Neurogenic differentiation (neurod) genes; Harold M. Weintraub, deceased, et al., 435/325, 69.1, 69.4, 172.3, 252.33, 320.1, 357, 360; 536/23.1, 23.5, 23.51 [IMAGE AVAILABLE]

US PAT NO: 5,695,995 [IMAGE AVAILABLE] L2: 1 of 10

ABSTRACT:

Neurogenic differentiation genes and proteins are identified, isolated, and sequenced. Expression of neuroD has been demonstrated in neural, pancreatic, and gastrointestinal cells. Ectopic expression of neuroD in non-neuronal cells of Xenopus embryos induced formation of neurons.

2. 5,693,487, Dec. 2, 1997, Nucleic acids encoding max: a helix-loop-helix zipper protein that forms a sequence-specific DNA-binding complex with Myc and Mad; Elizabeth M. Blackwood, et al., 435/69.1, 70.1, 172.3, 252.3, 320.1; 536/23.1, 23.5; 935/11, 22, 66, 70 [IMAGE AVAILABLE]

US PAT NO: 5,693,487 [IMAGE AVAILABLE] L2: 2 of 10

ABSTRACT:

Nucleic acid molecules capable of hybridizing under stringent conditions to the nucleotide sequence of the max cDNAs shown in SEQ ID NO: 1 or SEQ ID NO: 2, or to the nucleotide sequence of the mad cDNAs shown in SEQ ID NO: 5. The Max polypeptide when associated with the Myc or Mad

polypeptide is capable of binding to nucleotide sequences containing CACGTG.

3. 5,686,266, Nov. 11, 1997, Human brain sodium dependent inorganic phosphate cotransproter and related nucleic acid compounds; Binhui Ni, et al., 435/69.1, 7.1, 252.3, 320.1; 536/23.1 [IMAGE AVAILABLE]

US PAT NO:

5,686,266 [IMAGE AVAILABLE]

L2: 3 of 10

ABSTRACT:

This invention describes a novel human brain Na.sup.+ -dependent inorganic phosphate cotransporter, designated the hBNPI protein. This invention also encompasses nucleic acids encoding this protein, or a fragment thereof, as well as methods employing this protein and the nucleic acid compounds.

4. 5,652,224, Jul. 29, 1997, Methods and compositions for gene therapy for the treatment of defects in lipoprotein metabolism; James M. Wilson, et al., 514/44; 424/93.21; 435/172.3, 320.1, 325, 354, 366, 369, 370 [IMAGE AVAILABLE]

US PAT NO:

5,652,224 [IMAGE AVAILABLE]

L2: 4 of 10

ABSTRACT:

The invention provides a recombinant viral vector comprising the DNA of, or corresponding to, at least a portion of the genome of an adenovirus, which portion is capable of infecting a hepatic cell; and a human VLDL receptor gene operatively linked to regulatory sequences directing its expression. The vector is capable of expressing the normal VLDL receptor gene product in hepatic cells in vivo or in vitro. This viral vector is useful in the treatment of metabolic disorders caused by the accumulation of LDL in plasma, such as familial hypercholesterolemia or familial combined hyperlipidemia.

5. 5,624,818, Apr. 29, 1997, Nucleic acids encoding regulatory proteins that dimerize with Mad or Max; Robert N. Eisenman, et al., 435/69.1, 70.1, 172.3, 252.3, 320.1; 536/23.1, 23.5; 935/11, 22, 70, 72 [IMAGE AVAILABLE]

US PAT NO:

5,624,818 [IMAGE AVAILABLE]

L2: 5 of 10

ABSTRACT:

An isolated nucleic acid molecule capable of hybridizing under stringent conditions to the mSinA nucleotide sequence shown in FIG. 22 (SEQ ID NO:11), the mSin9A nucleotide sequence shown in FIG. 28 (SEQ ID NO:17), and/or the mSinB nucleotide sequence shown in FIG. 30 (SEQ ID NO:19). This isolated nucleic acid molecule preferably encodes a recombinant polypeptide which associates with a Mad polypeptide to form a recombinant polypeptide:Mad complex, which preferably associates with a Max polypeptide to form a recombinant polypeptide:Mad:Max complex, which preferably binds to a nucleotide sequence comprising CACGTG (SEQ ID NO:16). An isolated nucleic acid molecule capable of hybridizing under stringent conditions to a nucleotide sequence selected from among clone 10 shown in FIG. 24 (SEQ ID NO:9), clone 18 shown in FIG. 25 (SEQ ID NO:10), clone 19 shown in FIG. 26 (SEQ ID NO:11), and clone 20 shown in FIG. 27 (SEQ ID NO:12). This isolated nucleic acid molecule preferably encodes a recombinant polypeptide capable of associating with a Max polypeptide.

6. 5,618,918, Apr. 8, 1997, Human brain sodium dependent inorganic phosphate cotransporter; Binhui Ni, et al., 530/350, 300 [IMAGE AVAILABLE]

US PAT NO:

5,618,918 [IMAGE AVAILABLE]

L2: 6 of 10

ABSTRACT:

This invention descripe a novel human brain Na.sup.+ pendent inorganic phosphate cotransporter, designated the hBNPL protein. This invention also encompasses nucleic acids encoding this protein, or a fragment thereof, as well as methods employing this protein and the nucleic acid compounds.

7. 5,618,677, Apr. 8, 1997, Human brain sodium dependent inorganic phosphate cotransporter assay; Binhui Ni, et al., 435/7.1, 7.2, 69.1, 252.3, 320.1 [IMAGE AVAILABLE]

US PAT NO: 5,618,677 [IMAGE AVAILABLE]

L2: 7 of 10

ABSTRACT:

This invention describes a novel human brain Na.sup.+ -dependent inorganic phosphate cotransporter, designated the hBNPI protein. This invention also encompasses nucleic acids encoding this protein, or a fragment thereof, as well as methods employing this protein and the nucleic acid compounds.

8. 5,580,564, Dec. 3, 1996, Method for modifying the cell, tissue or host tropism of microorganisms; recombinant microorganisms obtained in this way and use thereof in medicine and veterinary medicine; Koenraad L. Glazenburg, et al., 424/229.1, 815; 435/172.1, 172.3, 236; 536/23.1, 24.1 [IMAGE AVAILABLE]

US PAT NO: 5,580,564 [IMAGE AVAILABLE] L2: 8 of 10

ABSTRACT:

The invention relates to a microorganism having a modified cell, tissue or host tropism whereby at least one gene of the microorganism, preferably an essential gene, is brought under the control of a nucleotide sequence specific for the cell, the tissue or the host. The specific nucleotide sequence can be a **promoter** sequence and/or enhancer sequence, which can be **inducible**. The invention is also directed at the use of such a recombinant microorganism for the provision of protection against the corresponding natural microorganism.

9. 5,552,309, Sep. 3, 1996, Use of polyols for improving the introduction of genetic material into cells; Keith L. March, 435/172.3; 424/93.1, 93.2, 426; 435/235.1, 320.1; 514/44; 935/57 [IMAGE AVAILABLE]

US PAT NO: 5,552,309 [IMAGE AVAILABLE] L2: 9 of 10

ABSTRACT:

A process for introducing an expression vehicle (e.g., plasmids, retroviral vectors, adenoviral vectors) into cells, which comprises contacting the cells with the expression vehicle and a polyol. The polyol may be a polyoxalkylene block copolymer, such as a polyoxypropylene-polyoxyethylene block copolymer. The use of the polyol provides for greater efficiency of transduction of the expression vehicle.

10. 5,302,519, Apr. 12, 1994, Method of producing a Mad polypeptide; Elizabeth M. Blackwood, et al., 435/69.1, 6, 69.3, 70.21; 530/350, 351; 536/23.1, 23.5 [IMAGE AVAILABLE]

US PAT NO: 5,302,519 [IMAGE AVAILABLE] L2: 10 of 10

ABSTRACT:

Nucleic acid molecules capable of hybridizing under stringent conditions to the nucleotide sequence residing between positions 1 and 453 of the max cDNAs shown in FIG. 2, or to the nucleotide sequence residing between positions 148 and 810 of the mad cDNAs shown in FIG. 14. The Max polypeptide when associated with the Myc or Mad polypeptide is capable of binding to nucleotide sequences containing CACGTG.

v.		

HILIGHT set on as ''
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Set Items Description
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? s adenovir? and E2A and promoter?
Processed 20 of 57 files ...
Processing
Processed 50 of 57 files ...
Completed processing all files
          125113 ADENOVIR?
            3297 E2A
          612970 PROMOTER?
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      s_3
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                       (Item 1 from file: 5)
DIALOG(R) File
              5:BIOSIS PREVIEWS(R)
(c) 1998 BIOSIS. All rts. reserv.
7682447
            BIOSIS Number: 90050447
  THE EMBRYONAL CARCINOMA STEM CELL ELA-LIKE ACTIVITY INVOLVES A
DIFFERENTIATION-REGULATED TRANSCRIPTION FACTOR
  LA THANGUE N B; THIMMAPPAYA B; RIGBY P W J
  LAB. EUKARYOTIC MOL. GENET., NATL. INST. MED. RES., THE RIDGEWAY, MILL
HILL, LONDON NW7 1AA, UK.
  NUCLEIC ACIDS RES 18 (10). 1990. 2929-2938.
                                                 CODEN: NARHA
  Full Journal Title: Nucleic Acids Research
  Language: ENGLISH
                                 - end of record -
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                       (Item 2 from file: 5)
      Display 3/3/2
DIALOG(R) File 5:BIOSIS PREVIEWS(R)
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6943828

43828 BIOSIS Number: 87004349
CYCLIC AMP INDUCTION OF EARLY ADENOVIRUS PROMOTERS IN LVES

SEQUENCES REQUIRED FOR E1A TRANS-ACTIVATION

SASSONE-CORSI P

MOL. BIOL. VIROL. LAB., SALK INST., P.O. BOX 85800, SAN DIEGO, CALIF. 92138.

PROC NATL ACAD SCI U S A 85 (19). 1988. 7192-7196. CODEN: PNASA Full Journal Title: Proceedings of the National Academy of Sciences of the United States of America

Language: ENGLISH

- end of record -

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Display 3/3/3 (Item 3 from file: 5) DIALOG(R) File 5:BIOSIS PREVIEWS(R) (c) 1998 BIOSIS. All rts. reserv.

6613502 BIOSIS Number: 86080053

AN ADENOVIRUS E1A-LIKE TRANSCRIPTION FACTOR IS REGULATED DURING THE DIFFERENTIATION OF MURINE EMBRYONAL CARCINOMA STEM CELLS

LA THANGUE N B; RIGBY P W J

LAB. EUKARYOTIC MOL. GENETICS, NATL. INST. MED. RES., THE RIDGEWAY, MILL HILL, LONDON NW7 1AA, UK.

CELL 49 (4). 1987. 507-514. CODEN: CELLB

Full Journal Title: Cell

Language: ENGLISH

- end of record -

Display 3/3/4 (Item 4 from file: 5) DIALOG(R) File 5:BIOSIS PREVIEWS(R) (c) 1998 BIOSIS. All rts. reserv.

BIOSIS Number: 85047193

A CELLULAR PROTEIN ACTIVATING TRANSCRIPTION FACTOR ACTIVATES TRANSCRIPTION OF MULTIPLE E1A-INDUCIBLE ADENOVIRUS EARLY

PROMOTERS

LEE K A W; HAI T-Y; SIVARAMAN L; THIMMAPPAYA B; HURST H C; JONES N C; GREEN M R

DEP. BIOCHEM. AND MOL. BIOL., HARVARD UNIV., 7 DIVINITY AVE., CAMBRIDGE, MASS. 02138.

PROC NATL ACAD SCI U S A 84 (23). 1987. 8355-8359. CODEN: PNASA Full Journal Title: Proceedings of the National Academy of Sciences of the United States of America

Language: ENGLISH

- end of record -

Display 3/3/5 (Item 1 from file: 154) DIALOG(R) File 154:MEDLINE(R) (c) format only 1998 The Dialog Corp. All rts. reserv.

04809110 86062906

Isolation and analysis of adenovirus type 5 mutants containing deletions in the gene encoding the DNA-binding protein.

Rice SA; Klessig DF

J Virol (UNITED STATES) Dec 1985, 56 (3) p767-78, ISSN 0022-538X Journal Code: KCV

Contract/Grant No.: AI 17315, AI, NIAID; T32GM07464-08, GM, NIGMS

Languages: ENGLISH

Document type: JOURNAL ARTICLE

?

English

(Item 1 from file: 351) Display 3/3/6 DIALOG(R) File 351: DERWENT WPI (c) 1998 Derwent Info Ltd. All rts. reserv. 010754580 **Image available** WPI Acc No: 96-251535/199625 XRAM Acc No: C96-079575 New replication-deficient adenoviral vectors having lethal early region gene deletions - useful in treatment of hereditary and acquired diseases, cancer gene therapies, and vaccines for prevention of infectious diseases Patent Assignee: CELL GENESYS INC (CELL-N) Inventor: FINER M H; JIA X; WANG Q Number of Countries: 067 Number of Patents: 003 Patent Family: Patent No Kind Date Applicat No Kind Date Week WO 9614061 A1 19960517 WO 95US14793 A 19951103 A61K-031/00 199625 B AU 9641092 A 19960531 WO 95US14793 A 19951103 A61K-031/00 199639 AU 9641092 A 19951103 EP 797436 A1 19971001 EP 95939149 A 19951103 A61K-031/00 199744 -more-? Display 3/3/6 (Item 1 from file: 351) DIALOG(R) File 351: DERWENT WPI (c) 1998 Derwent Info Ltd. All rts. reserv. WO 95US14793 A 19951103 Priority Applications (No Type Date): US 94333680 A 19941103 Filing Details: Patent Kind Filing Notes Application Patent WO 9614061 A1 Designated States (National): AL AM AU BB BG BR BY CA CN CZ EE FI GE HU IS JP KG KP KR KZ LK LR LS LT LV MD MG MK MN MX NO NZ PL RO RU SG SI SK TJ TM TT UA UZ VN Designated States (Regional): AT BE CH DE DK ES FR GB GR IE IT KE LS LU MC MW NL OA PT SD SE SZ UG WO 9614061 AU 9641092 A Based on EP 797436 Al Based on WO 9614061 Designated States (Regional): AT BE CH DE DK ES FR GB GR IE IT LI LU MC NL PT SE Language, Pages: WO 9614061 (E, 110); EP 797436 (E) - end of record -? (Item 1 from file: 399) Display 3/3/7 DIALOG(R) File 399:CA SEARCH(R) (c) 1998 American Chemical Society. All rts. reserv. 105147272 CA: 105(17)147272k **JOURNAL** Identification of a factor in HeLa cells specific for an upstream transcriptional control sequence of an EIA-inducible adenovirus promoter and its relative abundance in infected and uninfected cells AUTHOR(S): SivaRaman, Lakshmi; Subramanian, Subhalakshmi; Thimmappaya, Bayar LOCATION: Med. Sch., Northwestern Univ., Chicago, IL, 60611, USA JOURNAL: Proc. Natl. Acad. Sci. U. S. A. DATE: 1986 VOLUME: 83 NUMBER: 16 PAGES: 5914-18 CODEN: PNASA6 ISSN: 0027-8424 LANGUAGE:

?

Display 3/3/8 (Item 2 from file: 399) DIALOG(R) File 399:CA SEARCH(R) (c) 1998 American Chemical Society. All rts. reserv. CA: 104(19)162762t **JOURNAL** 104162762 Negative regulatory sequences in the EIa-inducible enhancer of the adenovirus-2 early EIIa promoter AUTHOR(S): Jalinot, P.; Kedinger, C. LOCATION: INSERM, Univ. Louis Pasteur, 67085, Strasbourg, Fr. JOURNAL: Nucleic Acids Res. DATE: 1986 VOLUME: 14 NUMBER: 6 PAGES: 2651-69 CODEN: NARHAD ISSN: 0305-1048 LANGUAGE: English - end of record -? Display 3/3/9 (Item 1 from file: 76) DIALOG(R) File 76: Life Sciences Collection (c) 1998 Cambridge Sci Abs. All rts. reserv. 01403997 2344693 The environment carcinoma stem cell Ela-like activity involves a differentiation-regulated transcription factor. Thangue, N.B.L.; Thimmappaya, B.; Rigby, P.W.J. Lab. Eukaryotic Mol. Genet., Natl. Inst. Med. Res., The Ridgeway, Mill Hill, London NW7 1AA, UK NUCLEIC ACIDS RES. vol. 18, no. 10, pp. 2929-2938 (1990.) DOCUMENT TYPE: Journal article LANGUAGE: ENGLISH SUBFILE: Biochemistry Abstracts Part 2: Nucleic Acids - end of record -? Display 3/3/10 (Item 2 from file: 76) DIALOG(R) File 76: Life Sciences Collection (c) 1998 Cambridge Sci Abs. All rts. reserv. 01203697 1891347 Cyclic AMP induction of early adenovirus promoters involves sequences required for E1A trans-activation. Sassone Corsi, P. Mol. Biol. and Virol. Lab., Salk Inst., P.O. Box 85800, San Diego, CA 92138, USA PROC. NATL. ACAD. SCI. USA. vol. 85, no. 19, pp. 7192-7196 (1988.) DOCUMENT TYPE: Journal article LANGUAGE: ENGLISH SUBFILE: Virology Abstracts; Genetics Abstracts; Biochemistry Abstracts Part 2: Nucleic Acids - end of record -(Item 1 from file: 149) Display 3/3/11 DIALOG(R) File 149: IAC(SM) Health & Wellness DB(SM) (c) 1998 Info Access Co. All rts. reserv. SUPPLIER NUMBER: 19553136 (USE FORMAT 7 OR 9 FOR FULL TEXT) Transcriptional regulation of the surfactant protein-A gene in fetal lung. (Thomas L. Petty 39th Annual Aspen Lung Conference: Genes and Gene Therapy) Mendelson, Carole R.; Gao, Erwei; Young, Pampee P.; Michael, Laura F.; Alcorn, Joseph L. Chest, v111, n6, p96S(9)

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June, 1997
PUBLICATION FORMAT: M zine/Journal ISSN: 0012-3692 ANGUAGE: English RECORD TYPE: Fulltext TARGET AUDIENCE: Professional
                     LINE COUNT: 00435
WORD COUNT:
              5298
                                  - end of record -
?
      Display 3/3/12
                         (Item 2 from file: 149)
DIALOG(R) File 149: IAC(SM) Health & Wellness DB(SM)
(c) 1998 Info Access Co. All rts. reserv.
                                           (USE FORMAT 7 OR 9 FOR FULL TEXT)
             SUPPLIER NUMBER: 04066237
01083741
Repression of the immunoglobulin heavy chain enhancer by the
  adenovirus-2 E1A products.
Hen, Rene; Borrelli, Emoliana; Chambon, P.
Science, v230, p1391(4)
Dec 20, 1985
PUBLICATION FORMAT: Magazine/Journal
                                      ISSN: 0036-8075 LANGUAGE: English
RECORD TYPE: Fulltext TARGET AUDIENCE: Academic
WORD COUNT:
             1835 LINE COUNT: 00180
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?
      Display 3/3/13
                         (Item 1 from file: 357)
DIALOG(R) File 357: Derwent Biotechnology Abs
(c) 1998 Derwent Publ Ltd. All rts. reserv.
                                        PATENT
199759 DBA Accession No.: 96-09939
New replication-deficient adenoviral vectors having lethal early
    region gene deletions - adeno virus and adeno-associated virus-based
    vector for human gene therapy and genetic immunization
AUTHOR: Wang Q; Finer M H; Jia X C
CORPORATE SOURCE: Foster City, CA, USA.
PATENT ASSIGNEE: Cell-GeneSys 1996
PATENT NUMBER: WO 9614061 PATENT DATE: 960517 WPI ACCESSION NO.:
    96-251535 (9625)
PRIORITY APPLIC. NO.: US 333680 APPLIC. DATE: 941103
NATIONAL APPLIC. NO.: WO 95US14793 APPLIC. DATE: 951103
LANGUAGE: English
                                  - end of record -
?
      Display 3/3/14
                         (Item 1 from file: 434)
DIALOG(R) File 434: Scisearch(R) Cited Ref Sci
(c) 1998 Inst for Sci Info. All rts. reserv.
           Genuine Article#: WM911 No. References: 37
15524709
Title: Conditional repression of the E2 transcription unit in E1-E3-deleted
    adenovirus vectors is correlated with a strong reduction in viral
    DNA replication and late gene expression in vitro
Author(s): Rittner K (REPRINT); Schultz H; Pavirani A; Mehtali M
Corporate Source: TRANSGENE SA, GENE THERAPY DEPT, 11 RUE MOLSHEIM/F-67000
    STRASBOURG//FRANCE/ (REPRINT)
Journal: JOURNAL OF VIROLOGY, 1997, V71, N4 (APR), P3307-3311
                 Publication date: 19970400
ISSN: 0022-538X
Publisher: AMER SOC MICROBIOLOGY, 1325 MASSACHUSETTS AVENUE, NW,
    WASHINGTON, DC 20005-4171
Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)
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(Item 2 from file: 434) Display 3/3/15 DISPLAY 3/3/15
DIALOG(R) File 434:Sci cch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. 12719632 Genuine Article#: MJ341 No. References: 68 Title: E2A EXPRESSION, NUCLEAR-LOCALIZATION, AND IN-VIVO FORMATION OF DNA-BINDING AND NON-DNA-BINDING SPECIES DURING B-CELL DEVELOPMENT Author(s): JACOBS Y; VIERRA C; NELSON C Corporate Source: UNIV CALIF RIVERSIDE, DEPT BIOCHEM/RIVERSIDE//CA/92521; UNIV CALIF RIVERSIDE, DEPT BIOCHEM/RIVERSIDE//CA/92521 Journal: MOLECULAR AND CELLULAR BIOLOGY, 1993, V13, N12 (DEC), P7321-7333 ISSN: 0270-7306 Document Type: ARTICLE (Abstract Available) Language: ENGLISH - end of record -? Display 3/3/16 (Item 3 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: LV644 No. References: 57 12545429 Title: THE E2A GENE-PRODUCT CONTAINS 2 SEPARABLE AND FUNCTIONALLY DISTINCT TRANSCRIPTION ACTIVATION DOMAINS Author(s): ARONHEIM A; SHIRAN R; ROSEN A; WALKER MD Corporate Source: WEIZMANN INST SCI, DEPT BIOCHEM/IL-76100 REHOVOT//ISRAEL/; WEIZMANN INST SCI, DEPT BIOCHEM/IL-76100 REHOVOT//ISRAEL/ Journal: PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA, 1993, V90, N17 (SEP 1), P8063-8067 ISSN: 0027-8424 Document Type: ARTICLE (Abstract Available) Language: ENGLISH - end of record -? Display 3/3/17 (Item 4 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: GX118 11274924 No. References: 69 Title: ANALYSIS OF VIRAL AND CELLULAR GENE-EXPRESSION DURING PROGRESSION AND SUPPRESSION OF THE TRANSFORMED PHENOTYPE IN TYPE 5 ADENOVIRUS -TRANSFORMED RAT EMBRYO CELLS Author(s): DUIGOU GJ; SU ZZ; BABISS LE; DRISCOLL B; FUNG YKT; FISHER PB Corporate Source: COLUMBIA UNIV COLL PHYS & SURG, INST CANC RES, DEPT PATHOL, CTR CANC, 650 W 168TH ST/NEW YORK//NY/10032; COLUMBIA UNIV COLL PHYS & SURG, INST CANC RES, DEPT PATHOL, CTR CANC, 650 W 168TH ST/NEW YORK//NY/10032; COLUMBIA UNIV COLL PHYS & SURG, INST CANC RES, DEPT UROL, CTR CANC/NEW YORK//NY/10032; COLUMBIA UNIV COLL PHYS & SURG, INST CANC RES, DEPT NEUROL SURG, CTR CANC/NEW YORK//NY/10032; ROCKEFELLER UNIV/NEW YORK//NY/10021; CHILDRENS HOSP, DEPT OPHTHALMOL/LOS ANGELES//CA/90054; UNIV SO CALIF/LOS ANGELES//CA/90054 Journal: ONCOGENE, 1991, V6, N10 (OCT), P1813-1824 Language: ENGLISH Document Type: ARTICLE (Abstract Available) - end of record -?

Display 3/3/18 (Item 5 from file: 434) DIALOG(R)File 434:Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv.

10812073 Genuine Article#: FH326 No. References: 24
Title: INTERFERENCE WITH PROTEIN-BINDING AT AP2 SITES BY SEQUENCE-SPECIFIC
METHYLATION IN THE LATE E2A PROMOTER OF ADENOVIRUS

TYPE-2 DNA Author(s): HERMANN R; Author(s): HERMANN R; ERFLER W
Corporate Source: UNIVOLOGNE, INST GENET, WEYERTAL 121) 5000 COLOGNE 41//FED REP GER/; UNIV COLOGNE, INST GENET, WEYERTAL 121/D-5000 COLOGNE 41//FED REP GER/ Journal: FEBS LETTERS, 1991, V281, N1-2, P191-195 Language: ENGLISH Document Type: ARTICLE (Abstract Available) - end of record -? Display 3/3/19 (Item 6 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. 10685884 Genuine Article#: EZ331 No. References: 36 Title: MULTICOMPONENT DIFFERENTIATION-REGULATED TRANSCRIPTION FACTORS IN F9-EMBRYONAL CARCINOMA STEM-CELLS Author(s): SHIVJI MKK; LATHANGUE NB Corporate Source: NATL INST MED RES, EUKARYOT MOLEC GENET LAB, RIDGEWAY MILL HILL/LONDON NW7 1AA//ENGLAND/; NATL INST MED RES, EUKARYOT MOLEC GENET LAB, RIDGEWAY MILL HILL/LONDON NW7 1AA//ENGLAND/ Journal: MOLECULAR AND CELLULAR BIOLOGY, 1991, V11, N3, P1686-1695 (Abstract Available) Language: ENGLISH Document Type: ARTICLE - end of display ->>>Page beyond end of display invalid ? d s3/9/15 (Item 2 from file: 434) Display 3/9/15 DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. Genuine Article#: MJ341 Number of References: 68 12719632 Title: E2A EXPRESSION, NUCLEAR-LOCALIZATION, AND IN-VIVO FORMATION OF DNA-BINDING AND NON-DNA-BINDING SPECIES DURING B-CELL DEVELOPMENT Author(s): JACOBS Y; VIERRA C; NELSON C Corporate Source: UNIV CALIF RIVERSIDE, DEPT BIOCHEM/RIVERSIDE//CA/92521; UNIV CALIF RIVERSIDE, DEPT BIOCHEM/RIVERSIDE//CA/92521 Journal: MOLECULAR AND CELLULAR BIOLOGY, 1993, V13, N12 (DEC), P7321-7333 ISSN: 0270-7306 Document Type: ARTICLE Language: ENGLISH Geographic Location: USA Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences Journal Subject Category: BIOCHEMISTRY & MOLECULAR BIOLOGY Abstract: A monoclonal antibody (Yae) was characterized and shown to specifically recognize E2A proteins in vivo, including the E2A-Pbx1 fusion gene products, p77E2A-Pbx1 and p85E2A-Pbx1. E2A proteins of a predominant molecular mass of 72 kDa, which -more-? Display 3/9/15 (Item 2 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv. comigrated with in vitro-produced rat E12 and rat E47, were detected in human pro-B, pre-B, mature B, and plasma cell lines. The Yae antibody detected an E2A-containing muE2 enhancer element-binding complex

(BCF-1) in pre-B- and mature B-cell lines in electrophoretic mobility shift assays which displayed a migration rate similar to that of in

vitro-produced rat E12 and rat E47. A new E2A-containing

muE2-binding species (P-E2A) was identified in plasma cells by

using electrophoretic mobility shift assays. **E2A** proteins were detected in pro-Balls but were unable to bind the uE2 site. These observations suggest that the muE2 site is the target of stage-specific **E2A** regulatory complexes during B-cell development. Immunostaining analyses demonstrated the predominant nuclear localization of **E2A** proteins. Finally, we have identified an **E2A** form, designated I-**E2A**, which is unable to bind DNA. Our observations demonstrate novel in vivo mechanisms for the regulation of transcription by **E2A** proteins during B-cell development.

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Display 3/9/19 (Item 6 from file: 434) DIALOG(R) File 434: Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv.

10685884 Genuine Article#: EZ331 Number of References: 36
Title: MULTICOMPONENT DIFFERENTIATION-REGULATED TRANSCRIPTION FACTORS IN
F9-EMBRYONAL CARCINOMA STEM-CELLS

Author(s): SHIVJI MKK; LATHANGUE NB

Corporate Source: NATL INST MED RES, EUKARYOT MOLEC GENET LAB, RIDGEWAY MILL HILL/LONDON NW7 1AA//ENGLAND/; NATL INST MED RES, EUKARYOT MOLEC GENET LAB, RIDGEWAY MILL HILL/LONDON NW7 1AA//ENGLAND/

Journal: MOLECULAR AND CELLULAR BIOLOGY, 1991, V11, N3, P1686-1695

Language: ENGLISH Document Type: ARTICLE

Geographic Location: ENGLAND

Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences
Journal Subject Category: BIOCHEMISTRY & MOLECULAR BIOLOGY
Abstract: Murine F9 embryonal carcinoma (F9 EC) stem cells have an Ela-like
transcription activity that is down-regulated as these cells

transcription activity that is down-regulated as these cells differentiate to parietal endoderm. For the adenovirus E2A promoter, this activity requires at least two sequence-specific

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transcription factors, one that binds the cyclic AMP-responsive element (CRE) and the other, DRTF1, the DNA-binding activity of which is down-regulated as F9 EC cells differentiate. Here we report the characterization of several binding activities in F9 EC cell extracts, referred to as DRTF 1a, 1b and 1c, that recognize the DRTF1 cis-regulatory sequence (-70 to -50 region). These activities can be chromatographically separated but are not distinguishable by DNA sequence specificity. Activity la is a detergent-sensitive complex in which DNA binding is regulated by phosphorylation. In contrast, activities 1b and 1c are unaffected by these treatments but exist as multicomponent protein complexes even before DNA binding. Two sets of DNA-binding polypeptides, p50DR and p30DR, affinity purified from F9 EC cell extracts produce complexes 1b and 1c. Both polypeptides appear to be present in the same DNA-bound protein complex and both directly contact DNA. These affinity-purified polypeptides activate transcription in vitro in a binding-site-dependent manner. These data indicate the in F9 EC stem cells, multicomponent

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Display 3/9/19 (Item 6 from file: 434) DIALOG(R)File 434:Scisearch(R) Cited Ref Sci (c) 1998 Inst for Sci Info. All rts. reserv.

differentiation-regulated transcription factors contribute to the cellular Ela-like tivity.

cellular Ela-like tivity.

Identifiers--KeyWords Flus: DEPENDENT ACTIVATION; BINDLE PROTEINS; TRANS-ACTIVATION; ElA GENE; PROMOTER; EXPRESSION; DNA; E2F; EMBRYOGENESIS; PURIFICATION

Research Fronts: 89-1013 003 (SEQUENCE-SPECIFIC DNA INTERACTION OF THE FOS JUN PROTEIN COMPLEX; EARLY GENE INDUCTION; TRANSCRIPTION FACTOR AP-1; LEUCINE ZIPPER DOMAIN)

89-3178 002 (LIVER-SPECIFIC PYRUVATE-KINASE GENE **PROMOTER**; NUCLEAR FACTOR-I; INVITRO TRANSCRIPTION; **ADENOVIRUS** DNA-BINDING PROTEIN; CCAAT BOX SEQUENCE)

89-4793 002 (IMMUNOGLOBULIN HEAVY-CHAIN **PROMOTER**; INVITRO TRANSCRIPTION; C-MYC GENE; OCTAMER MOTIF; **INDUCIBLE** NUCLEAR FACTORS; UPSTREAM MUSCLE-SPECIFIC ENHANCER)

89-7387 001 (EMBRYONAL CARCINOMA-CELLS; RETINOIC ACID-INDUCED NEURAL DIFFERENTIATION; TESTIS-SPECIFIC ANTIGEN OF THE C57BL/6 MOUSE)
Cited References:

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Display 3/9/10 (Item 2 from file: 76) DIALOG(R)File 76:Life Sciences Collection (c) 1998 Cambridge Sci Abs. All rts. reserv.

01203697 1891347

Cyclic AMP induction of early **adenovirus promoters** involves sequences required for E1A trans-activation.

Sassone Corsi, P.

Mol. Biol. and Virol. Lab., Salk Inst., P.O. Box 85800, San Diego, CA 92138, USA

PROC. NATL. ACAD. SCI. USA. vol. 85, no. 19, pp. 7192-7196 (1988.)

DOCUMENT TYPE: Journal article LANGUAGE: ENGLISH

SUBFILE: Virology Abstracts; Genetics Abstracts; Biochemistry Abstracts Part 2: Nucleic Acids

Early in adenovirus infection, the E1A (early region 1A) oncogene products trans-activate the other early viral transcription units, as well as some cellular promoters. The mechanism by which E1A elicits its activity is still unknown. In this report, the authors show that adenovirus E2a and E3 promoters are cAMP inducible

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Display 3/9/10 (Item 2 from file: 76)
DIALOG(R)File 76:Life Sciences Collection
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in rat pheochromocytoma PC12 cells and that this activation requires the presence of the cAMP-dependent protein kinase II. Using deletion mutants of the E2a promoter, it was found that the sequence TACGTCAT located between positions -70 and -77 is involved in both the cAMP response and the E1A trans-activation.

DESCRIPTORS: adenovirus; promoters; cyclic AMP; protein kinase; gene expression

IDENTIFIERS: early region; induction; rats; pheochromocytoma cells; effects on; trans-activation

SECTION HEADING: 22044 --Viral nucleic acid synthesis & synthesis of virus-coded proteins; 07313 --Viruses; 14662 --Gene regulation

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